# Analysis of Transmission Dynamics of Anthrax in Animal Population. A modeling Approach

#### 4 Abstract

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5 This paper seeks to develop a SIR model with vaccination compartment in the study of 6 anthrax transmission dynamics in animal population. The model employ ordinary differential 7 equations in the formulation of the model's equation. The model's steady states solutions are 8 investigated. The disease free equilibrium and endemic equilibrium of the model are analyzed 9 qualitatively. Vaccination rate below a certain critical value causes the anthrax disease to 10 persist. Recruitment and contact rates are the most sensitive parameters that contribute significantly to the basic reproductive ratio. 11 

#### 12 **1.0 Introduction**

Anthrax is an infectious disease categorized under zoonotic diseases caused by a bacterium 13 14 called Bacillus anthracis [4]. The disease is found naturally in soil [19] and mostly affects 15 herbivores. Anthrax is one of the major diseases that cause uncontrolled mortality in cattle, 16 pigs, sheep, goats and horses worldwide [3, 10, 15]. Animals easily get infected with anthrax 17 through contact with infected animals, consumption of infected grass or water and by 18 inhalation of anthrax spores [18]. The environment is usually infected with carcasses from 19 infected animals. Grass and soil become the reservoirs of anthrax spores which can persist in 20 the soil or grass for an extended period of time even under very extreme weather and 21 environmental conditions.

22 Authors [1] model consists of susceptible, contamination, infective and pathogens. The model 23 regards infective compartment as key to the transmission of anthrax. In this model, the 24 infected animals do show clinical signs of the disease. According to author [2], the model 25 consists of susceptible, contamination and pathogens. The model does not include infective 26 compartment. According to the model, the infective compartment do have very low 27 reproductive ratio [11] and does not cause any infections in animals.

28 Research done by authors [8] the model considers transmission, carcass ingestion, 29 environment and migration as possible means through which anthrax is transmitted in 30 animals. In this model, carcass ingestion and removal of carcasses from the environment does 31 not cause any decline of anthrax transmission in animals.

### 32 **2.0 The Model**

This model divides the total animal population at any time (t) into four sub compartments with respect to their disease status in the system. The total animal population is given by N (t) =S(t)+I(t)+R(t)+V(t) where S (t) represents animals at risk of developing anthrax infection, I (t) represents animals showing anthrax symptoms, R (t) represents animals recovered from anthrax infection and acquired temporal immunity and V (t) represents animals susceptible and are vaccinated against anthrax attack.

39 The parameters used in this model are:  $\lambda$  denotes recruitment rate;  $\beta$  denotes contact rate;  $\mu$ 40 denotes natural death rate;  $\gamma$  denotes vaccination rate;  $\tau$  denotes waning immunity of 41 vaccinated animals;  $\sigma$  denotes waning recovery rate;  $\theta$  denotes disease induced death rate 42 and  $\alpha$  animal recovery rate.

The diagram below shows SIR model flow chart with vaccination compartment for anthraxtransmission in animal population.

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Figure 1: SIR Flow chart with vaccination compartment

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49 The model equations are:

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$$\frac{dS}{dt} = \lambda - \beta SI - (\mu + \gamma)S + \sigma R + \tau V$$

$$\frac{dI}{dt} = \beta SI - (\mu + \theta + \alpha)I$$

$$\frac{dR}{dt} = \alpha I - (\mu + \sigma)R$$

$$\frac{dV}{dt} = \gamma S - (\mu + \tau)V$$
(1)

51 Disease Free Equilibrium is given by  $\varepsilon^0 = (S^0, I^0, R^0, V^0)$ . There exists no anthrax disease 52 and no animals are infected with anthrax. The critical point is given by  $\varepsilon^0 = (\frac{\lambda}{\mu + \gamma}, 0, 0, 0)$ .

According to authors [3, 11], reproductive ratio can be found using Jacobian matrix J of (1)as:

55 J (SIRV) = 
$$\begin{pmatrix} -(\mu+\gamma) & -\beta\frac{\lambda}{\mu+\gamma} & \sigma & \tau \\ 0 & \beta\frac{\lambda}{\mu+\gamma} - (\mu+\theta+\alpha) & 0 & 0 \\ 0 & \alpha & -(\mu+\sigma) & 0 \\ 0 & 0 & 0 & -(\mu+\tau) \end{pmatrix}$$
56 (2)  
57 Determinant of (2) become  $1 - \beta \frac{\lambda}{(\mu+\gamma)(\mu+\theta+\alpha)}$   
58 (3)  
59  $\beta \frac{\lambda}{(\mu+\gamma)(\mu+\theta+\alpha)}$   
60 (4) Expression (4) is called the basic reproductive  
61 ratio  $R_0$ .  
62 **Theorem 1**

63 Disease free equilibrium point is locally asymptotically stable if  $R_0 < 1$  and anthrax disease 64 will not persist. 65 If  $R_0 > 1$ , disease free equilibrium become unstable.

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67  
68  
69 **Proof**  
70 Disease free equilibrium point 
$$\varepsilon^{*}(\frac{\lambda}{\mu+\gamma}, 0, 0, 0)$$
 has reproductive ratio given  
71  $R_{0} = \beta \frac{\lambda}{\mu+\gamma} - (\mu+\theta+\alpha)$   
72 At disease free equilibrium  $\beta \frac{\lambda}{\mu+\gamma} - (\mu+\theta+\alpha) < 0$   
73 (5)  
74 Equation (5) can be expressed as  $\beta \frac{\lambda}{(\mu+\gamma)(\mu+\theta+\alpha)} - 1 < 0$ .  
75  $R_{0} = \beta \frac{\lambda}{(\mu+\gamma)(\mu+\theta+\alpha)}$ 

as

Therefore, 
$$R_0 - 1 < 0$$
 which implies that  $R_0 < 1$ .

Given that  $R_0 < 1$ , we have disease free equilibrium point which is locally asymptotically stable.

## 79 Lemma 1

80 If 
$$\beta \frac{\lambda}{(\mu+\gamma)} - (\mu+\theta+\alpha) > 0$$
, then it follows that  $\beta \frac{\lambda}{(\mu+\gamma)(\mu+\theta+\alpha)} - 1 > 0$ 

- 81 Therefore,  $R_0 > 1$  which implies that the disease will persist in animal population.
- According to authors in [15], endemic equilibrium of dynamical system (1) is given by  $\varepsilon^* = (S^*, I, R^*, V^*)$  where  $S^* > 0, I^* > 0, R^* > 0$  and  $V^* > 0$ .
- 84 From (2), the rest point becomes:

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$$=\frac{\mu+\theta+\alpha}{\beta}$$

 $S^*$ 

86 
$$I^* = \frac{(\mu+\gamma)(\mu+\tau)(\mu+\theta+\alpha) - \gamma\tau(\mu+\theta+\alpha) - \beta\lambda(\mu+\sigma)}{\beta(\mu+\tau)[\sigma\tau - (\mu+\sigma)(\mu+\theta+\alpha)]}(\mu+\sigma)$$

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$$R^* = \tau \frac{(\mu + \gamma)(\mu + \tau)(\mu + \theta + \alpha) - \gamma \tau(\mu + \theta + \alpha) - \beta \lambda(\mu + \tau)}{\beta(\mu + \tau) [\sigma \tau - (\mu + \sigma)(\mu + \theta + \alpha)]} , \quad V^* = \gamma \frac{(\mu + \theta + \alpha)}{\beta(\mu + \tau)}$$

- 88 (6)
- 89 If the vaccination is less than a certain critical value, the disease persist.
- 90
- 91
- 92
- 93 Theorem 2
- 94 If  $\gamma < \frac{\beta\lambda}{(\mu + \theta + \alpha)} \mu$ , then the endemic equilibrium point become unstable. The disease will
- 95 persist.
- 96 **Proof**

97 From the Jacobian matrix 
$$\mathbf{J} = \begin{pmatrix} -(\mu + \gamma) & -\frac{\beta\lambda}{\mu + \gamma} & \sigma & \tau \\ 0 & \beta\frac{\lambda}{\mu + \gamma} - (\mu + \theta + \alpha) & 0 & 0 \\ 0 & \alpha & -(\mu + \sigma) & 0 \\ 0 & 0 & 0 & -(\mu + \tau) \end{pmatrix}$$

98 The determinant is greater than zero

99 
$$\frac{\beta\lambda}{(\mu+\gamma)} - (\mu+\theta+\alpha) > 0$$
 (7)

100 Re-arranging (7) yields

101 
$$\frac{\beta\lambda}{(\mu+\gamma)(\mu+\theta+\alpha)} - 1 > 0$$
(8)

102 But from (4), 
$$R_0 = \frac{\beta\lambda}{(\mu + \gamma)(\mu + \theta + \alpha)}$$

103 Therefore, 
$$R_0 > 1$$
.

104 The endemic equilibrium will only occur if  $R_0 > 1$ . This means that the disease become 105 unstable and the rest point is lost. The vaccinated animals loose their immunity and become 106 susceptible.

(9)

107 Lemma 2

108 If  $\gamma > \frac{\beta\lambda}{\mu + \theta + \alpha} - \mu$ , the endemic equilibrium point becomes stable. Therefore, the disease

109 persists.

110 The table below shows sensitivity analysis on how each parameter contribute to the basic 111 reproductive ratio  $R_0$  of the model. Sensitivity analysis is given by the relation:

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$$S_A^{R_0} = \frac{\partial R_0}{\partial A} \times \frac{A}{R_0}$$

114 Where A is any parameter used in the model.

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Parameter	Contribution	Baseline values	References
λ	Positive	200	[4]
β	Positive	0.0001	[1,7]
μ	Negative	0.001	Estimate
γ	Negative	0.10	[7]
σ	Negative	0.02	Estimate
τ	Negative	0.003	[3]

θ	Negative	0.15	[1]
α	Negative	0.01	[7]

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118 **3.0 Results and Discussion** 

In this study, we modeled vaccination compartment in the transmission dynamics of anthrax
in animal population. The outcome of stability analysis of the endemic equilibrium state
shows that it is possible to effectively control anthrax outbreak in animal population.

Taking the initial conditions for endemic equilibrium  $\varepsilon^*$ ( $S^* = 2000, I^* = 100, R^* = 300, V^* = 500$ ) and time t= 10 years and considering parameters baseline values from other literature as indicated above, the reproductive ratio  $R_o = 1.2299$ . Increasing the rate of vaccination  $\gamma$ , the reproductive ratio  $R_o$  decreases. Therefore, animals will not die as a result of anthrax infection. When  $\gamma$  is increased by 24.5%, the reproductive ratio  $R_o$  decrease by 19.50%. In this case, reproductive ratio become 0.9900 which is less than unit. Hence, the disease free equilibrium.

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The outcome of the model shows that vaccination is a good control strategy against anthrax outbreak in animal population. However, vaccination may not completely guarantee protection of the animals against anthrax but it is possible that the vaccinated animals with time may lose immunity and may contract anthrax disease again. Therefore, there is need to keep vaccinating animals periodically against anthrax to keep anthrax prevalence as low as possible or completely eradicated.

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### 138 **<u>REFERENCES</u>**

139 Anderson, R, M., May, R, M (1979). Population biology of infectious diseases nature 280.

140 Bradford Gutting et al (2016).Mathematically modeling Inhalational anthrax

7

- Buddhi,P et al (2016).Optimal control applied in an anthrax epizootic model. Journal of
  biological Systems 24(04), p 495-517.
- 143 Castillo-Chavez C, et al (2002).Computation of  $R_o$  and its role in global stability. In:
- 144 Castillo-Chavez C et al (Eds) Mathematical approaches for emerging and re-emerging
- 145 Infectious diseases: an introduction, vol125.IMA, p 229-250.
- 146 Friedman, A., Yakubu, A, A (2013). Anthrax epizootic and migration: persistence or
- extinction. Mathematics Bioscience volume 241, p 137-144.
- Hahn, B.D., Furniss, P, R (1983). A mathematical model of anthrax epizootic in the
  Kruger National Park. Applied mathematics model 5:130 model of anthrax epizootic
  threshold results.
- Hahn, B, D., Furniss, P, R (1979). A deterministic model of anthrax epizootic threshold
  results. Ecological modeling volume 20 issue 2-3, p 233-241.
- Kalu, A, Ugwa. Agwu, I, A & Akuagwu, A, N (2013) Mathematical Analysis of the
  Endemic Equilibrium of the transmission Dynamics of Tuberculosis vol. 2issue 12
- LaSalle, J.S. (1976). The stability of dynamical systems. CBMS-NSF regional conference
  series in applied mathematics, volume 25.SIAM, Philadelphia.
- Mushayabasa, S. (2015). Global stability of an anthrax model with Environmental
  decontamination and time delay. Discrete Dynamics in Nature and Society, Article ID
  573146.
- Mushayabasa, S. (2011). Impact of vaccination and culling on controlling foot and mouth
  disease: A mathematical modeling approach. World J Vaccines 1, p 156-161.
- Osman, S., Oluwole, D. M., & Theuri, D. M. (2018). Mathematical model of transmission
  dynamics of anthrax in human and animal population, vol.8 no 6.
- Pauline van den D, Watmough, J. (2002). Reproduction Numbers and sub-threshold
  endemic equilibria for compartmental models of disease transmission. Journal
  Mathematical Biosciences volume 180, issue 1-2, p 29-48

167	Saad-Roy, C, M., van den Pauline., Yakubu, A, A (2017).Mathematical model of anthrax
168	transmission in animal populations. Bulletin of Mathematical Biology, vol.79 no.2 pp303-
169	324
170	Sudipa, C., Misra O, P., Dhar, J (2014). Stability Analysis of SIR model with vaccination
171	p17-23
172	Wendy, C, T (2013).Soil ingestion, nutrition and seasonality of anthrax in herbivores of
173	Etosha National Park, Ecosphere 4(1), p1-19
174	World Health Organization (2008). Anthrax in humans and animals, 4th edition, World
175	Health Organization p10.
176	World Health Organization (2016). Anthrax in Humans and animals. Geneva:
177	International Office of Epizootics, World Health Organization
178	Yusuf, T, T., Benyah, F (2012) Optimal control of vaccination and treatment for an SIR
179	epidemiological model. World Journal of modeling and Simulation, vol. 8 no.3 pp194-
180	204
181	Zerihun, M.S., Narasimha, M.S (2016). Modeling and Simulation Study of Anthrax
182	Attack on Environment vol. 3 issue 4.Journal of multidisciplinary Engineering and
183	Technology (JMEST)
184	
185	
186	
187	
188	
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